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## STATUS OF THE CLAIMS:

## 1.-47. (Canceled)

- 48. (Previously Presented) A method for providing analgesia in a subject, said method comprising delivering a composition comprising fentanyl or a fentanyl congener to the subject, wherein the composition is administered to the subject using an implantable convective delivery system, is delivered from the system for 48 hours or more at a low volume rate of 2 ml/day or less and is sufficient to provide analgesia in the subject.
- 49. (Previously Presented) The method of claim 48, wherein the composition is delivered using a patterned delivery regime.
- 50. (Previously Presented) The method of claim 49, wherein the composition is delivered in a substantially continuous fashion.
- 51. (Previously Presented) The method of claim 49, wherein the composition is delivered in a substantially uninterrupted manner for a pre-selected period of time.
- 52. (Previously Presented) The method of claim 49, wherein the composition is delivered in a substantially constant fashion.
- 53. (Previously Presented) The method of claim 49, wherein the composition is delivered over an extended period of time.
- 54. (Previously Presented) The method of claim 53, wherein the composition is delivered for a period of about 72 hours.

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55. (Previously Presented) The method of claim 53, wherein the composition is delivered for a period from 2 to 5 days.

- 56. (Previously Presented) The method of claim 53, wherein the composition is delivered for a period of at least about 100 days.
- 57. (Previously Presented) The method of claim 49, wherein the composition is delivered using a controlled drug delivery device.
- 58. (Previously Presented) The method of claim of claim 57, wherein the controlled delivery device is implanted in the subject's body.
- 59. (Previously Presented) The method of claim 48, wherein the composition is delivered to the subject at a volume rate of from about 0.01 µl/day to about 100 µl/day.
- 60. (Previously Presented) The method of claim 48, wherein the composition is delivered to the subject at a volume rate of from about 0.04 µl/day to about 10 µl/day.
- 61. (Previously Presented) The method of claim 48, wherein the composition is delivered to the subject at a volume rate of from about 0.2 µl/day to about 5 µl/day.
- 62. (Previously Presented) The method of claim 48, wherein the composition is delivered to the subject at a volume rate of from about 0.5 µl/day to about 1 µl/day.
- 63. (Previously Presented) A method for providing analgesia in a subject, said method comprising delivering to the subject a composition comprising fentanyl or a fentanyl congener, wherein said fentanyl or fentanyl congener is present in the composition at a concentration of about 0.5 mg/ml to about 500 mg/ml or greater, and further wherein the composition is administered to the subject using an implantable convective delivery system, is delivered from

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the system at a low volume rate of about 2 ml/day or less and is sufficient to provide analgesia in the subject.

- 64. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is in solution.
- 65. (Previously Presented) The method of claim 64, wherein the fentanyl or fentanyl congener is dissolved in a liquid carrier.
- 66. (Previously Presented) The method of claim 63, wherein the composition is administered to the subject as a semi-solid, gel, liquid, suspension, emulsion or an osmotic dosage pharmaceutical formulation.
- 67. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of at least about 2 to at least about 10,000 times greater than the solubility of fentanyl or fentanyl congener in aqueous solution.
- 68. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of from about 0.5 mg/ml to about 500 mg/ml.
- 69. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of from about 1 mg/ml to about 400 mg/ml.
- 70. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of from about 50 mg/ml to about 400 mg/ml.

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71. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of from about 75 mg/ml to about 300 mg/ml.

- 72. (Previously Presented) The method of claim 63, wherein the fentanyl or fentanyl congener is present in the composition at a concentration of from about 100 mg/ml to about 250 mg/ml.
- 73. (Previously Presented) The method of claim 63, wherein the composition is delivered at a low volume rate of 2 ml/day or less.
- 74. (Previously Presented) The method of claim 63, wherein the composition is delivered using a patterned delivery regime.
- 75. (Previously Presented) The method of claim 74, wherein the composition is delivered in a substantially continuous fashion.
- 76. (Previously Presented) The method of claim 74, wherein the composition is delivered in a substantially uninterrupted manner for a pre-selected period of time.
- 77. (Previously Presented) The method of claim 74, wherein the composition is delivered in a substantially constant fashion.
- 78. (Previously Presented) The method of claim 74, wherein the composition is delivered over an extended period of time.
- 79. (Previously Presented) The method of claim 78, wherein the composition is delivered for a period from about 2 to about 48 hours.

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80. (Previously Presented) The method of claim 78, wherein the composition is delivered for a period from about 2 to 5 days.

- 81. (Previously Presented) The method of claim 78, wherein the composition is delivered for a period of at least about 100 days.
- 82. (Previously Presented) The method of claim 74, wherein the composition is delivered using a controlled drug delivery device.
- 83. (Previously Presented) The method of claim of claim 82, wherein the controlled delivery device is implanted in the subject's body.
- 84. (Previously Presented) A method for providing analgesia in a subject, said method comprising delivering to the subject a composition comprising fentanyl or a fentanyl congener, wherein the composition is administered to the subject using an implantable convective delivery system, the composition is delivered from the system for 48 hours or more at a low volume rate sufficient to deliver from about 0.01 µg/hour to about 200 µg/hour of the fentanyl or fentanyl congener to the subject, and further wherein said amount of delivered fentanyl or fentanyl congener is sufficient to establish a systemic analgesic effect in the subject.
- 85. (Previously Presented) The method of claim 84, wherein the fentanyl or fentanyl congener is in solution.
- 86. (Previously Presented) The method of claim 85, wherein the fentanyl or fentanyl congener is dissolved in a liquid carrier.
- 87. (Previously Presented) The method of claim 84, wherein the composition is administered to the subject as a semi-solid, gel, liquid, suspension, emulsion or an osmotic dosage pharmaceutical formulation.

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88. (Previously Presented) The method of claim 84, wherein the systemic analgesic effect is sufficient to manage pain in the subject.

- 89. (Previously Presented) The method of claim 84, wherein the systemic analgesic effect is sufficient to treat pain in the subject.
- 90. (Previously Presented) The method of claim 84, wherein the systemic analgesic effect is sufficient to modulate pain response in the subject.
- 91. (Previously Presented) The method of claim 84, wherein the systemic analgesic effect is sufficient to ameliorate or alleviate pain in the subject.
- 92. (Previously Presented) The method of claim 48, wherein the fentanyl congener is sufentanil.
- 93. (Previously Presented) The method of claim 63, wherein the fentanyl congener is sufentanil.
- 94. (Previously Presented) The method of claim 84, wherein the fentanyl congener is sufentanil.
- 95. (Previously Presented) The method of claim 68, wherein the fentanyl congener is sufentanil.
- 96. (Previously Presented) The method of claim 69, wherein the fentanyl congener is sufentanil.

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97. (Previously Presented) The method of claim 70, wherein the fentanyl congener is sufentanil.

- 98. (Previously Presented) The method of claim 71, wherein the fentanyl congener is sufentanil.
- 99. (Previously Presented) The method of claim 72, wherein the fentanyl congener is sufentanil.